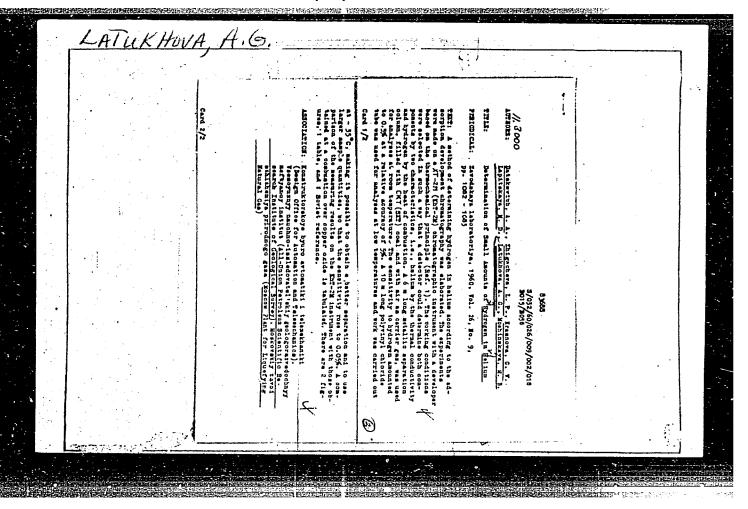


TURKEL'TAUB, N.M.; SHCHVARTSMAN, V.P.; KANCHEYEVA, O.A.; LATUKHOVA, A.G.; KOLYUBYAKINA, A.I.

Use of thermodynamic apparatus in gas surveys. Trudy VNIGHI no.11: 260-272 '58. (MIRA 13:1) (Gases--Analysis) (Geochemical prospecting)



34542 S/659/61/007/000/030/044 D217/D303

1.1800

AUTHORS: Gorbunov, N.S., Kovalev, Ye.A., and Latukhova, A.G.

TITLE:

Investigating diffusion coatings resistant to media

containing vanadium pentoxide

SOURCE:

Akademiya nauk SSSR. Institut metallurgii. Issledovaniya po zharoprochnym splavam, v. 7, 1961, 263 - 270

TEXT: In this investigation, in which the service conditions of gas transport turbines were simulated, the excess pressure of the working process and the speed of gas flow were not allowed for. The work was carried out at the Institut fizicheskoy khimii AN SSSR (Institute of Physical Chemistry AS USSR) and at the Vsesoyuznyy nauchno-issledovatel skiy institut zheleznodorozhnogo transporta (All Union Scientific Research Institute of Railway Transportation) in association with the Kolomenskiy teplovozostroitel nyy zavod im. Kuybyshev (Kolomensk Internal Combustion Works im. Kuybyshev). Diffusion coatings were produced on the surface of the austenitic class chromium-nickel steel 3N 417 (EI417), from which flat speci-

Card 1/4

X

Investigating diffusion coatings ...

S/659/61/007/000/030/044 D217/D303

mens, 15 x 10 x 6 mm were made. Silicide diffusion coatings were produced at 1000, 1020 and 1050°C by soaking for 2 - 6 hours. Aluminal and chromiding in vacuum at 1000°C for 4 - 6 hours. The force of the mode as the temperature for corrosion testing, this being the malected as the temperature for guide vanes of a gas turbine. To semens, at which the rate of corrosion of the specimens at elevated alloys in the course of service of the gas turbine plant, two meder and application of a suspension to the specimens in molten cintatures (painting). On testing the above coatings in an atmosphere of air in contact with cinder (10 and 41.6 % V<sub>2</sub>O<sub>5</sub>) at 730°C, silicided specimens was lower. All coatings, apart from silicided 730°C. The corrosive medium diffused through the coating to the me-

Investigating diffusion coatings ...

S/659/61/007/000/030/044 D217/D303

tal, oxidizing the latter at the boundary line of diffusion. The thickness of a silicided layer under similar conditions decreased somewhat and pitting corrosion appeared on the surface; however, molten cinder did not penetrate to the metal and the latter did not corrode. In the presence of SiO<sub>2</sub> in air atmosphere, the rate of

corrosion of alumino-silicided and aluminized specimens is the same as the rate of corrosion in pure air. Chromided and silicided specimens exhibit high stability under these conditions. A combination cementation coating (Si and Al) gave less protection to the steel EI417 against vanadium pentoxide than a coating consisting of one of the individual elements. On periodically cooling the specimens (cooling 40 times from 730 to 20°C within 15-20 minutes), no exfoliation and destruction of the protective layer of chromided and silicided specimens occurred. No cracks or ruptures in the diffusion layer were observed on water quenching silicided specimens from 1150°C and the adhesion of the coating to the base metal remained unimpaired. Siliciding and chromiding are recommended for protection of gas vanes of gas turbine plants against corrosion during combustion of sulphur-containing petroleum residues of high Card 3/4

X

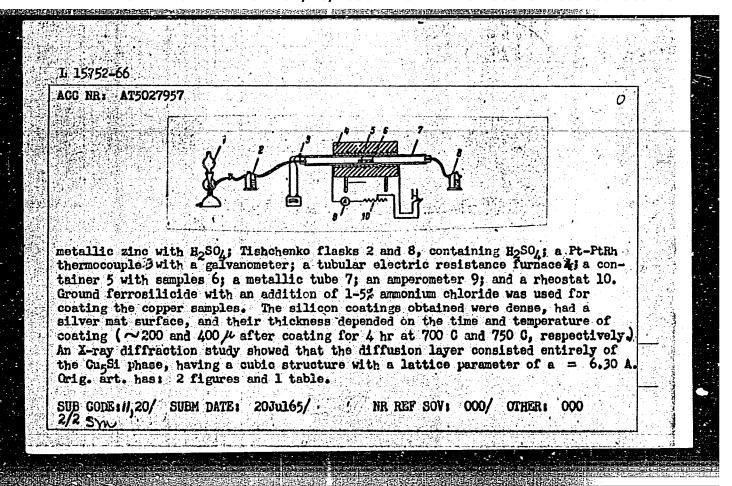
Investigating diffusion coatings ...

S/659/61/007/000/030/044 D217/D303

vanadium content. There are 5 figures, 2 tables and 11 references: 3 Soviet-bloc and 8 non-Soviet-bloc. The 4 most recent references to the English-language publications read as follows: Corrosion, 11 no. 1, p. 35, 1955; Iron and Steel Inst., 179, no. 4, p. 342, 1955; Corrosion, 12, no. 9, pp. 49-54, 1956; Iron and Steel Inst., 182, no. 2, p. 195, 1956.

Card 4/4

at the control of the first of the control of the c	SOURCE CODE: UR/0000/65/000/000/0216/0218
AUTHOR: Gorbunev. S. A.; Pa	N. S. (Doctor of chemical sciences); Latukhova, A. G.; Klevtsur, vloya, V. A.
ORG: Inone	60
TITLE: Diffusion	of silicon coatings on copper
SOURCE: Seminar p	oo zharostoykim pokrytiyam. Leningrad, 1964. Zharostoykiye
pokrytiya	(Heat-resistant coatings); trudy seminara. Leningrad, 12d-vo 265, 216-218
	trolyte, copper, silicon, internal stress, crystal lattice
stresses during ab	olytically applied coatings on copper suffered large internal brupt variations of temperature. This resulted in the cracking of the coatings. Experiments on the diffusion coating of copper.
Alak acmilan yana	made in the flow of dried hydrogen in an apparatus (see fig.) ipp generator 1 for the production of H by the reaction of
consisting of a Ki	



3 (9)

AUTHOR:

Latun, V. S.

SOV/50-59-8-8/19

TITLE:

Calculation of Level in Wind Tides and Wind Back Tides in the Taganrogskiy Gulf (Raschet urovnya pri sgonakh i nagonakh v

Taganrogskom zalive)

PERIODICAL:

Meteorologiya i gidrologiya, 1959, Nr 8, pp 29 - 31 (USSR)

ABSTRACT:

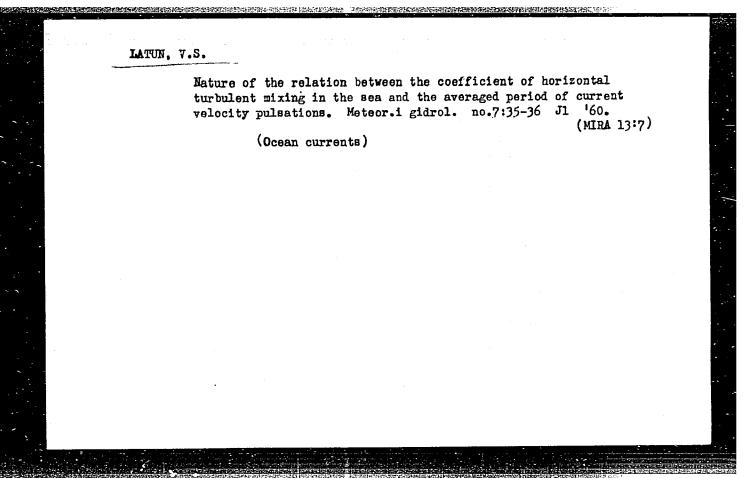
The Taganrogskiy Gulf has a length of 120 km, a width of 32 km and a depth of 6 m. The stretched form of the shallow-water gulf permits the calculation of the wind tides in this gulf to be regarded as a plane task of hydrodynamics. The level variations are expressed in such a case by ordinary differential equations with constant coefficients. The equation (1) of B. A. Tareyev (Ref 2), and the equation (2) of G. I. Gershengorn, are used for this purpose. Both are similar in structure, but not identical. Tareyev investigates the turbulent friction in the liquid, Gershengorn the friction on the bottom. Calculation formulas are derived from these two equations, and the results of calculations carried out by them are compared. The equations (1) and (2) are only applicable to closed basins. At the open end of the Taganrogskiy Gulf, an amplitude of level variations amounting to only one-tenth of those in the upper

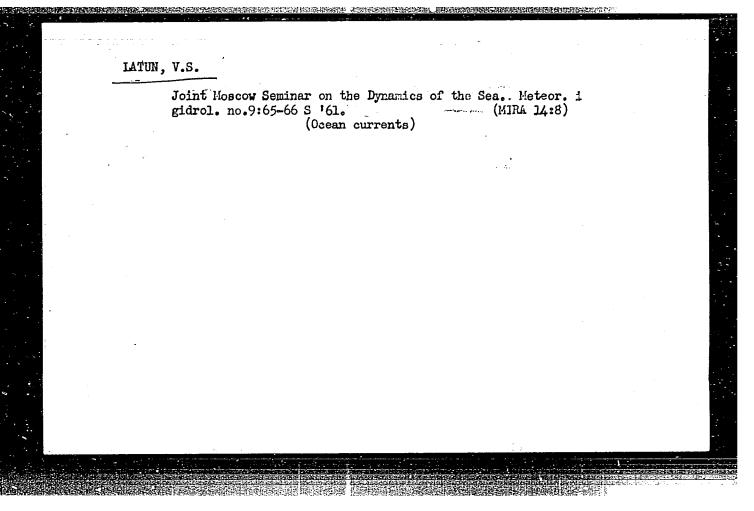
Card 1/2

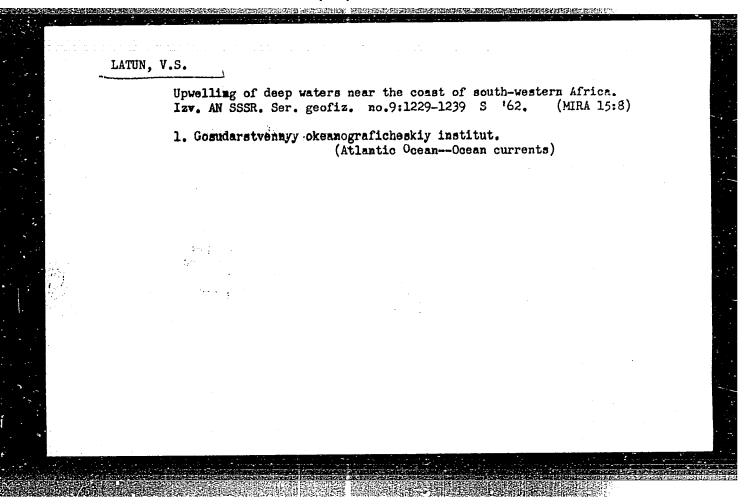
Calculation of Level in Wind Tides and Wind Back Tides in the Taganrogskiy Gulf sov/50-59-8-8/19

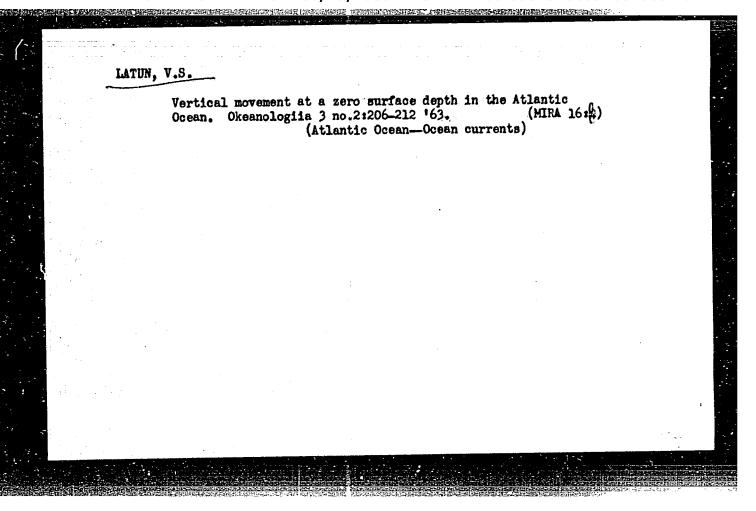
part of the gulf, are observed at wind tides and wind back tides. Formula (5) is derived from formula (2), and formulas (6a), (6b) and (6v) are derived from (1). 7 observations of wind tides and wind back tides are used to check these formulas. The results calculated by the formulas derived are put forward in figures 1 and 2, partly in form of a diagram. With their help it is shown that formula (5) yields a result nearly equal to reality, while among the formulas (6) the formula (6b) proves to be the best one. The calculations put forward allow the conclusion that the coefficient m of the variation damping with  $m = 0.12 \frac{1}{hour}$ , and the kinematic coefficient  $\nu$  of the horizontal turbulent exchange with  $\nu = 9.4 \cdot 10^8$  cm<sup>2</sup>/sec, are characteristic of the Taganrogskiy Gulf. There are 2 figures and 2 Soviet references.

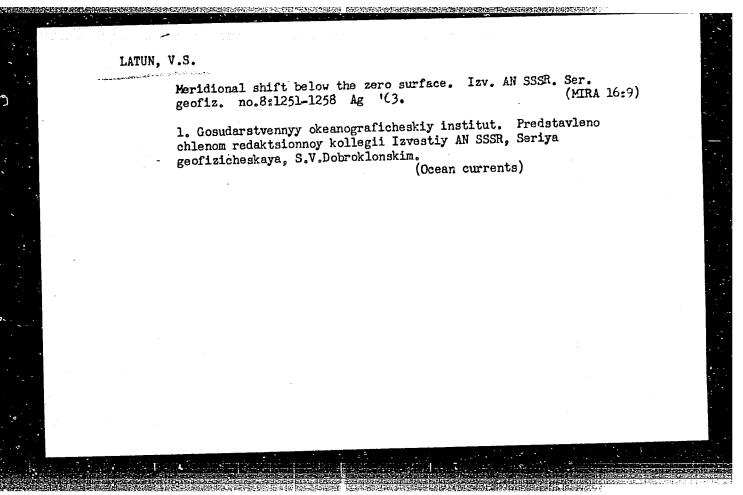
Card 2/2

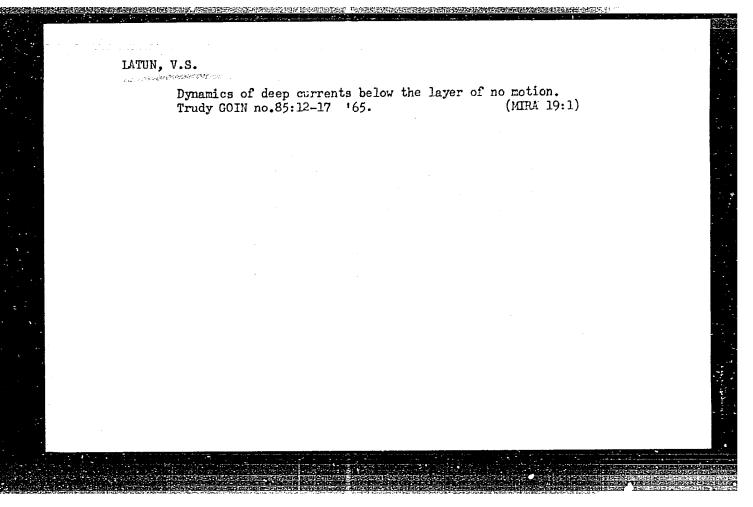


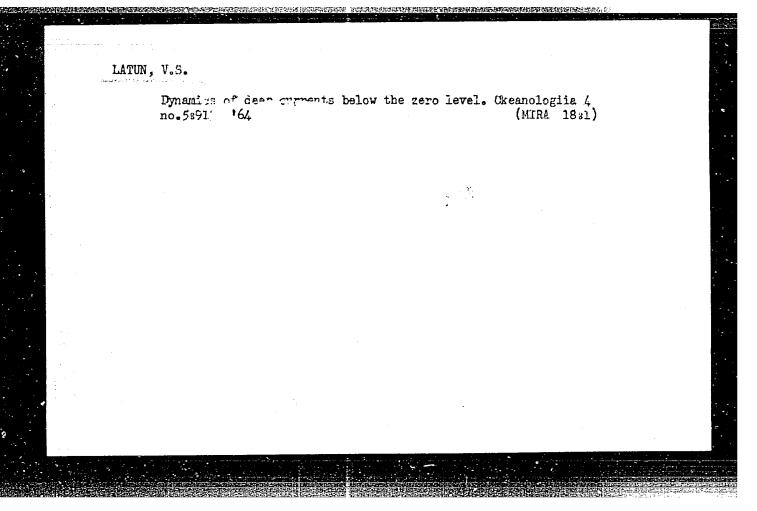












L 35997-66 EWT(1) GW (N)

SOURCE CODE: UR/2634/65/000/085/0012/0017

AUTHOR: Latun, V. S.

36 31

ORG: None

TITLE: The dynamics of deep currents below the zero level

SOURCE: Moscow. Gosudarstvennyy okeanograficheskiy institut. Trudy, no. 85, 1965. Teoriya i metody raschetov techeniy i neperiodicheskikh kolebaniy urovnya i prilivov (Theory and methods of calculating currents and acyclic fluctuations of water level and tides), 12-17

TOPIC TAGS: ocean current, ocean dynamics, fluid flow, fluid velocity

ABSTRACT: This paper presents improvements of the original model published earlier by the author (Okeanologiya, no. 2, 1963). The earlier boundary conditions, imposed on densities at z=0 and z= depth of the channel D, are now replaced by the new velocity condition v=0 at the bottom (z=D). The newly obtained solutions for densities and velocities, presented in the paper, are compared with the experimental data on deep sea currents in the Gulf Stream region and the western North Atlantic Ocean and are found in good qualitative

Card 1/2

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HILLINITY BENESHEVICH, I.I., kandidat tekhnicheskikh nauk; BOOIN, N.H., kandidat tekhnicheskikh nauk; BYKOV, Ye.i., inzhener; VLASOV, I.I., kandidat tekhmicheskikh nauk; GRITSEVSKIY, M.Ye., inzhener; GRUBER, L.O., inzhener; GURVICH, V.G., inzhener; DAVYDOV, V.H., inzhener; YER-SHOV, I.M., kandidat tekhnicheskikh nauk: ZASORIN, S.N., kandidat tekhnicheskikh neuk; IVANOV, I.I., kandidat tekhnicheskikh neuk; KRAUKLIS, A.A., inzhoner; ERUTOV, L.B., inzhener; LAPIN, V.B., inzhener; LASTOVSKIY, V.P., dotscut; LATUNIN, H.I., inzhener; MARKVAHDT, K.G., professor, doktor tekhnicheskikh nauk; MAKHAYLOV, M.I., professor, doktor tekhnicheskikh nauk; NIKANOROV, V.A., inzhener; OSKOLKOV, K.N., inzhener; OKHOSHIN, L.I., inzhener; PARFENOV, K.A., dotsent, kandidat tekhnicheskikh nauk; PERTSOVSKIY, L.M., inzhener; POPOV, I.P., inzhener; PGRSHNEV, B.G., inzhener; RATNER, H.P., inzhener; ROSSIYAVSKIY, G.I., dotsent, kandidat tekhnicheskikh nauk; RYKOY, I.I., kendidet tekhnicheskikh nauk; RYSHKOYSKIY, I.Ya., dotsent, kandidat teknnicheskikh nauk; RYABKOV, A.Ya., professor [deceased]: TAGER, S.A., kandidet tekhnicheskikh nauk: KHAZEN, M.M., professor, doktor tekhnicheskikh nauk; CHERNYSHEV, M.A., doktor teknnicheskikh nauk; HBiN, L.Ye., professor, doktor tekhnicheskikh nauk; TUKEMAY, B.M., dotsent; AKSKNOY, I.Ya., dotsent, kandidat tekhnicheskikh neuk; ARKHANGEL SKIT, A.S., inzhener; BARTENEV, P.V., professor, dekter teknnicheskikh nauk; EERNGARD, K.A., kandidat tekhnicheskikh nauk; BUNOVOT, N.Ye., dotsent, kandidat tekhnicheskikh nauk; BOSDANOV, I.a., inchener; BOSDANOV, N.K., kandidat tekhnicheskikh nauk; VINNICIMNEO, N.G., dotsent, kandidat ekonomicheskikh nauk; (Continued on next card)

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HENESHEVICH, I.I. --- (continued) Card 2.

VASIL'YEV, V.F.; GONCHAROV, N.G., inzhener; DERIBAS, A.T., inzhener; DOBROSEL'SKIY, K.H., dotsent, kandidat tekhnicheskikh nauk; DIUGACH, B.A., kandidat tekhnicheskikh nauk; YKFIMOV, G.P., kandidat tekhnicheskikh nauk; ZEMBLINOV, S.V., professor, doktor tekhnicheskikh nauk; ZABZLLO, M.L., kandidat tekhnicheskikh nauk; IL'IN, K.P., kandidat tekhnicheskikh nauk: KARWINIKOV, A.D., kandidat tekhnicheskikh nauk; KAPLUH, F.Sh., inzhener; KANSHIN, M.D.; KOCHNEV, P.P., professor, doktor tekhnicheskikh nauk; KOGAH, L.A., kandidat tekhnicheskikh nauk; KUCHURIN, S.F., inzhener; LEVASHOV, A.D., inzhener; MAKSIHOVICH, B.M., dotsent, kandidat tekhnicheskikh nauk; MARTYNOV, M.S., inzhener; MEDRL', O.M., inzhener; NIKITIN, V.D., professor, kandidat tekhnicheskikh nauk; PADNYA, V.A., inzhener; PANTELEYEV, P.I., kandidat tekhnicheskikh nauk; FNTROV, A.P., professor, doktor tekhnicheskikh nauk; POVOROZHENKO, V.V., professor, doktor tekhnicheskikh nauk; PISKAREV, I.I., dotsent, kandidat tekhnicheskikh nauk; SERGEYEV, Ye.S., kandidat tekhnicheskikh neuk; SIMONOV, K.S., kandidat tekhnichekikh nauk; SIMANOVSKIY, M.A., inzhener; SUYAZOV, I.G., inzhener; TAIDAYEV, F.Ya., inzhener; TIKHONOV, K.K., kendidat tekhnicheskikh nauk; USHAKOV, N.Ya., inzhenr; USFENSKIY, V.K., inzhener; FEL\*DMAN, B.D., kandidat tekhnicheskikh nauk; FERAPONTOV, G.V., inzhener; KHOKHLOV, L.P., inzhenr; CHERNCHORDIK, G.I., professor, doktor tekhnicheskikh nauk; SHAMAYEV, H.F., inshener; SHAFIRKIN, B.I., inzhener; YAKUSHIN, S.I., inzhener; GRANOVSKIY, P.G., redaktor; TISHGHENKO, A.I., redaktor; ISAYEV, I.P., dotsent, kandidat tekhnicheskikh nauk, redsktor; KLIMOV, V.F., dotsent kandidat tekhnicheskikh (Continued on next card)

I.ATUNIN, Nikolay Ivg govich; OKHOSHIN, Leonid Ivanovich; BELYAYKV, I.A., inzh., red.; KHITROV, P.A., tekhn.red.

[Handbook for an electrician in railroad power engineering]
Spravochnik elektromontera energeticheskogo khoziaistva zheleznykh dorog. Moskva, Gos.transp.zhel-dor.izd-vo, 1959.

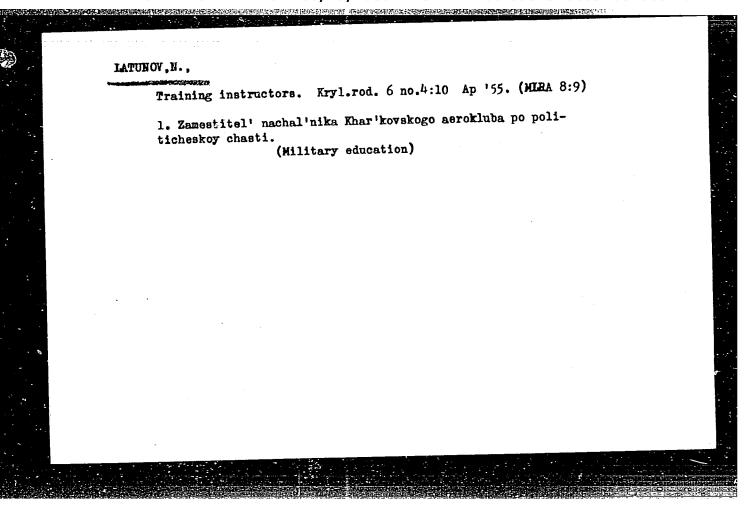
570 p. (MIRA 13:2)

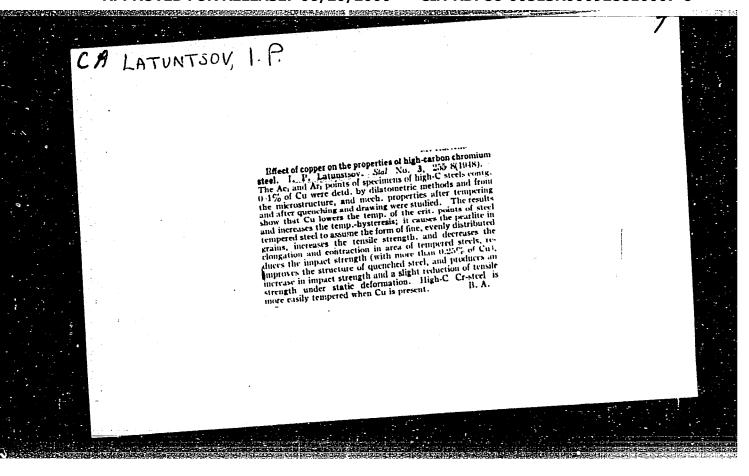
(Electric railroads)

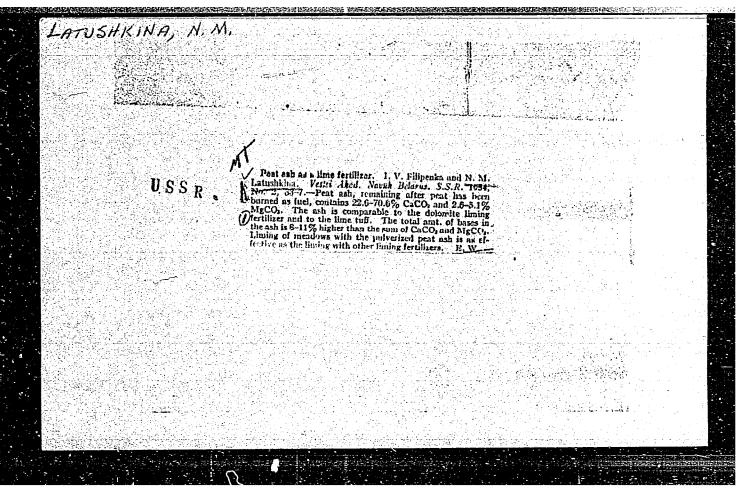
LATUNIN, Nikolay Ivanovich; OKHOSHIN, Leonid Ivanovich; ZATUCHNYY,
I.M., inzh., retsenzent; KALININ, V.K., kand. tekhn.nauk,
red.; USENKO, L.A., tekhn. red.

[Handbook for reilroad electric power
plants] Spravochnik elektromontera energeticheskogo khoziatstva zheleznykh dorog. Izd.2., perer. Moskva, Transzheldorizdat, 1963. 446 p.

(MIRA 17:2)







## LATUSHKINA, V. B.

LEVINA, A.I.; LATUSHKINA, V.B.

Comparative evaluation of the MIOT electroprecipitator (developed by the Moscow Research Institute for the Protection of Labor) and Green's sedimentator. Bor'ba s sil. 1:162-166 '53. (MLRA 7:10)

1. Moskovskiy nauchno-issledovatel'skiy institut okhrany truda VTsSPS.
(AIR--POLLUTION) (DUST)

LATUSHKINA, V. B.

Latushkina, V. B.

"Problems of labor hygiene in plants producing abrasive materials and abrasive tools (attempts to find a scientific absis for the maximum permissible content in the air of synthetic abrasive dust)." First Moscow Order of Lenin Medical Inst imeni I. M. Sechenov. Moscow, 1956. (Dissertation for the Degree of Candidate in Medical Sciences)

Knizhnaya letopis' No. 21, 1956. Moscow.

```
LATUSHKINA, V.B., nauchnyy sotrudnik

A new device for experimental research on the effect of industrial dust on the organism. Gig. i san. 21 no.8:18-24 Ag '56. (MIRA 9:11)

1. Iz Moskovskogo institutaokhrany truda VTSSPS i kafedry giglyeny truda I Moskovskogo ordena Lenina meditsinskogo instituta imeni I.M.Sechenova.

(INDUSTRIAL HYGIBUS

exper. research on eff. of indust. dust on organism)

(DUST

in indust., exper. research on eff.)
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LATUSHEIMA, V. F.

"Hygienic characteristics of the industrial dust of artificial abrasives."

report submitted at the 13th All-Union Congress of Hygienists, Epidemiologists and Infectionists, 1959.

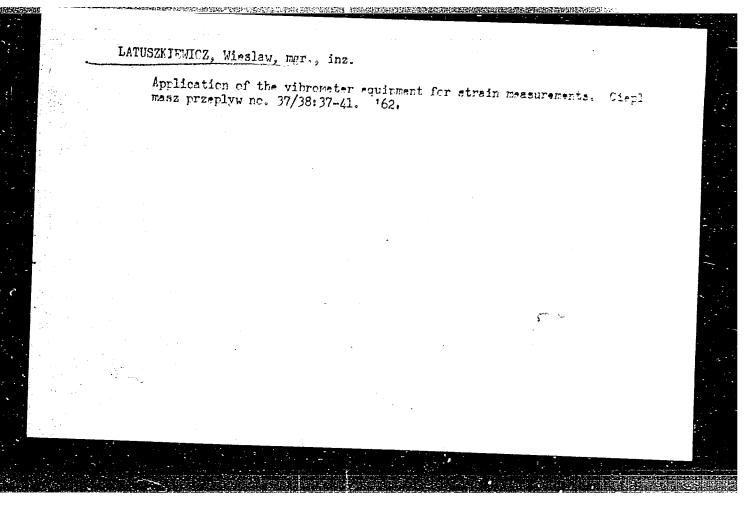
是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,这种是一个大学的,我们就是一个大学的,我们就是 第一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就是一个大学的,我们就

LATUSHKINA, V.B. (Moskva)

Changes in the respiratory organs under the action of dust from artifical abrasives. Gig. truda i prof. zab. 4 no. 7:49-52 '60. (MIRL 13:8)

1. Institut okhrany truda Vsesoyuznogo tsentral'nogo soveta profsoyuzov i kafedra gigiyeny truda I Moskovskogo ordena Lenina meditsinskogo instituta im. I.M. Sechenova.

(LUNGS--DUST DISEASES) (ABRASIVES--PHYSIOLOGICAL EFFECT)



62

ACCESSION NR: AT4025428

P/0000/62/000/000/0093/0099

AUTHOR: Latuszkiewicz, Wieslaw (Master engineer)

TITLE: Instrument for measurement of stresses and vibration frequency in rotating elements

SOURCE: Konferencja "Technika Pomiarowa w Cieplnych Maszynach Przeplywowych" (Conference on "Measurement technique in thermal flow machines"). Lodz, 1962, 93-99

TOPIC TAGS: stress measurement, turbine stress measurement, vibration frequency, vibration frequency measurement, turbine rotating element, NG-6 vibration measuring instrument

ABSTRACT: The NG-6 apparatus was developed and built by the Heat Engineering Institute's Department of Heat Turbines. This instrument is intended for measuring stresses and vibration frequencies in rotating elements. Measurements are by strain gauge method. The device is characterized by a high degree of simplicity and makes possible the use of one strain gauge in the rotating unit without the necessity of temperature compensation. The impulse from the strain

Card 1/3

# ACCESSION NR: AT4025428

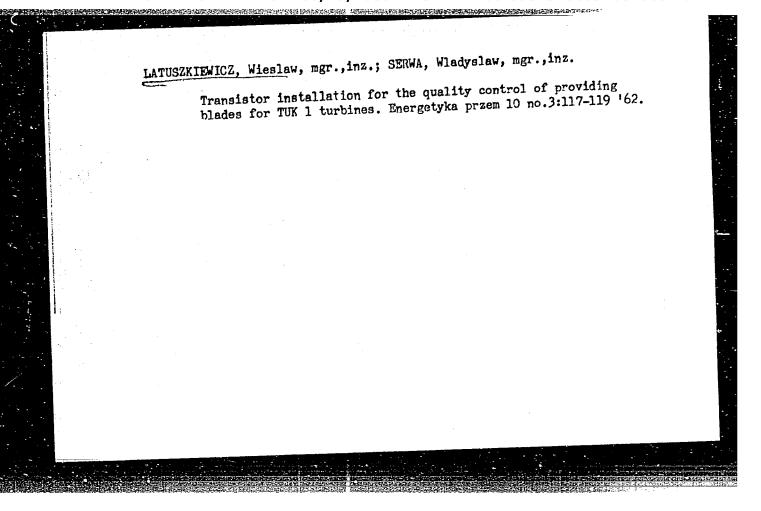
gauge is fed through a rotating relay and multipoint change-over switch into an amplifier. The amplifier is powered by a stabilized voltage feed. The amplified pulse is then transmitted to a bifilar suspension oscillograph or cathode oscilloscope. The scaler is connected in parallel to the working strain gauge at the multipoint relay's input. The operation of the apparatus is based on resistance strain gauges working in a circuit of resistance potentiometers. The amplifier has an amplification of 10°. Its frequency range is 20 to 10,000 kc and range of measured stresses is 50 to 4000 kg/cm² at a maximum output current of 100 milliamps. Any cathode type oscilloscope can be used to observe the pulses recorder can be hooked up to the amplifier's output for recording the responses at the amplifier output. If the vibration frequency of the tested element is of given frequency should be emitted and the measured frequency then determined by a comparison of signals. Orig. art. has: 3 figures.

ASSOCIATION: Instytut Techniki Cieplnej, Zaklad Turbin Cieplnych, Lods: (Heat Engineering Institute, Department of Heat Turbines)

Cord 2/3

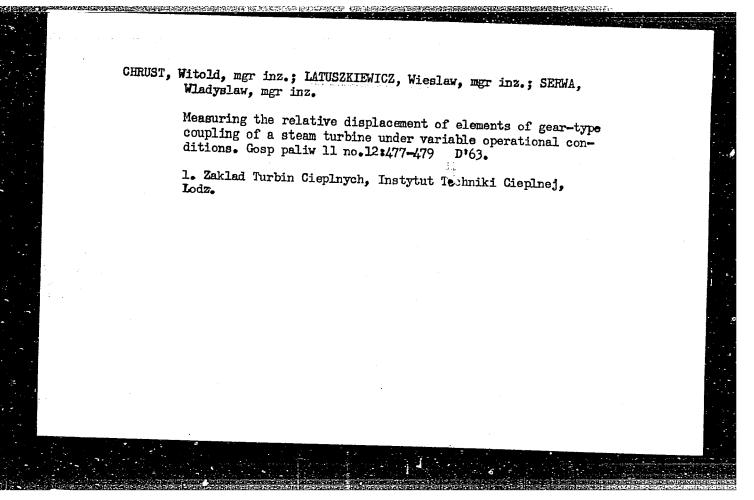
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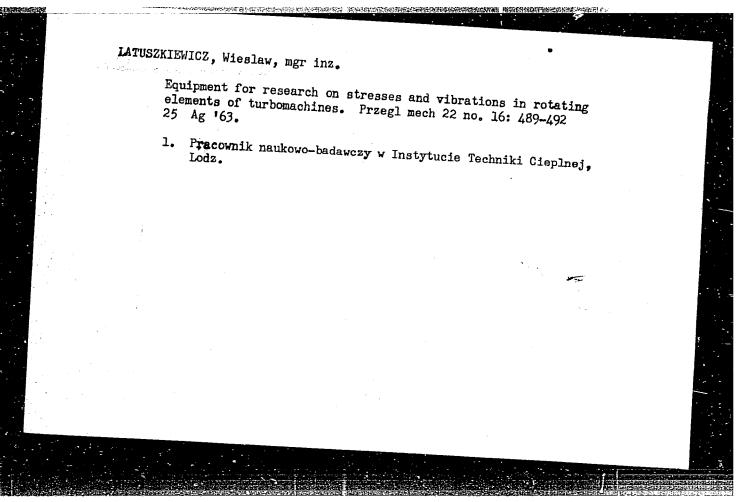
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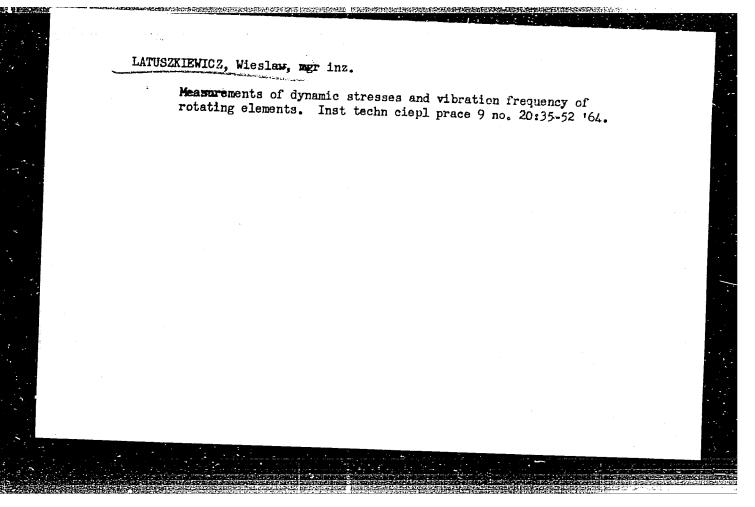


LATUSZKIEWICZ, Wieslaw, mgr inz.; LISICKI, Andrzej, mgr inz.

Testing of steam turbines for operational adjustment under conditions of deteriorated vacuum in the condenser. Energetyka przem 10 no.12:439-444 D '62.







LATUSZKIEWICZ, Wieslaw, mgr inz.; LISICKI, Andrzej, mgr inz.

Testing the 7 MW stal radial flow turbine in order to adjust it to operation under conditions of a worsened condenser vacuum. Gosp paliw 12 no.2:Suppl.: Biul inst techn ciepl 12 no.2:73-76 F '64.

1. Zaklad Turbin Cieplnych, Instytut Techniki Cieplnej, Lodz.

LATUSZKIEWICZ, Wieslaw, mgr inz.; LISICKI, Andrzej, mgr inz.

Tests of the 7 M/ steel radial-flow turbine in order to adjust it to work under conditions of worsened vacuum in the condenser. Biul inst techn ciepl 12 no.2:73-76 F '64.

1. Department of Thermal Turbines of the Institute of Heat Engineering, Lodz.

### LATVIAN, A.S.

The posibility of production of maleic anhydride and maleic dialdehyde from fur furol.

Report to be submitted for the 12th Conference on high molecular weight compounds devoted to monomers, Baku, 3-7- April 62

ATVIKOVA. P.S. U USSR/Physics - Oscillatory spectrum

FD-939

Card 1/1

Pub. 146 - 13/20

Author

: Latvikova, P. S. THE PROPERTY OF THE PARTY OF TH

Title

: Oscillatory structure in the spectra of zinc oxide

Periodical

: Zhur. eksp. i teor. fiz., 27, No 5 (11), 636-645, Nov 1954

Abstract

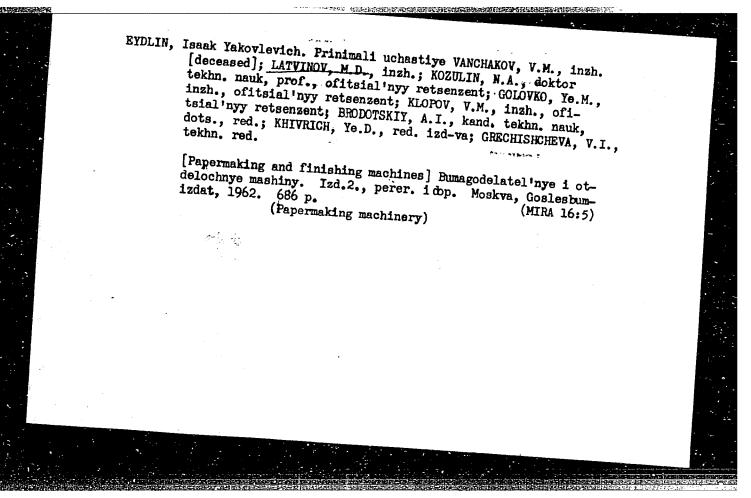
: In the absorption spectra and radiation spectra of zinc oxide one observes a system of weak maxima. The author describes the electronvibration formula for two values of the energy of electron perturbation. The electron transition is connected with the perturbation of the activator. In this case the oscillatory states of the lattice, which is connected energy-wise with the activator, appear in the spectra of absorption and radiation. Eleven references, 2 Western and 9 USSR (e.g. G. A. Konovalov, Dissertation, Tomsk State University, 1952. L. A. Suvolorova, Diploma Work, Tomsk State University, 1952. N. L. Gasting, Sib. FTI, Otchet za 1947 [Siberian Physico-technical Institute, Account for 1947].)

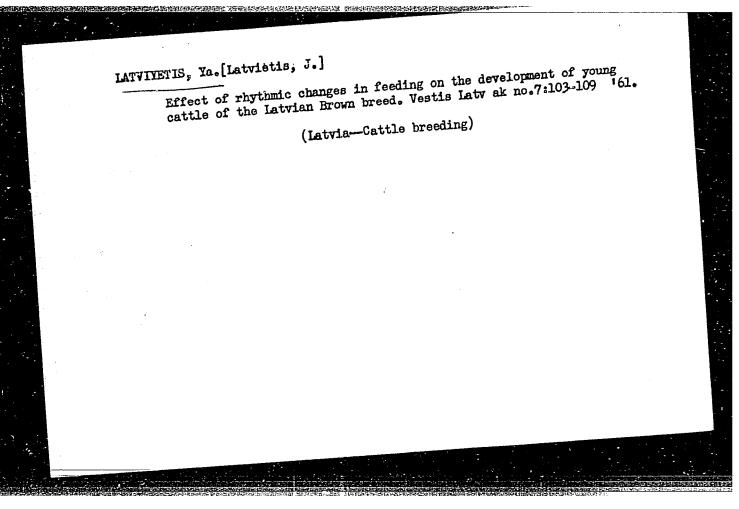
Institution : Siberian Physicotechnical Institute, Tomsk State University

Submitted

: November 18, 1953

CIA-RDP86-00513R000928810007-6" APPROVED FOR RELEASE: 06/20/2000

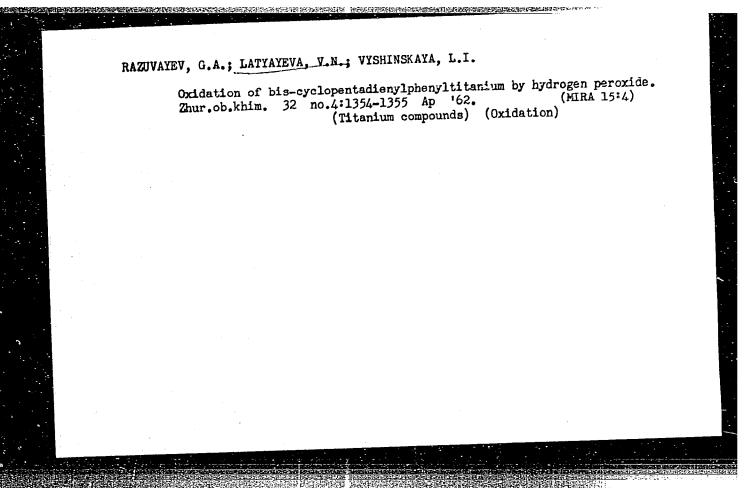




LATVYS, V.; SLIZYS, V.

Formation and determination of compounds in the system CaSO\_-SiO\_-Al\_2O\_3-Fe\_2O\_3-C. Trudy AN Lit. SSR. Ser.B no.l: 153-159 F62 (MIRA 17:8)

1. Institut khimii i khimicheskoy tekhnologii AN Litovskoy SSR.



LATYAYEVA, V.N.

USSE/Chemistry - Photoreaction

Pub. 151 - 14/38 Card 1/1

: Razuvaev, G. A.; Ol'dekop Yu. A.; and Latyaeva, V. N. Authors

Photoreaction of organometallic mercury compounds in solutions. Part 14.-Photoreaction of beta-mercuribispropionic acid and its dimethyl ether Title

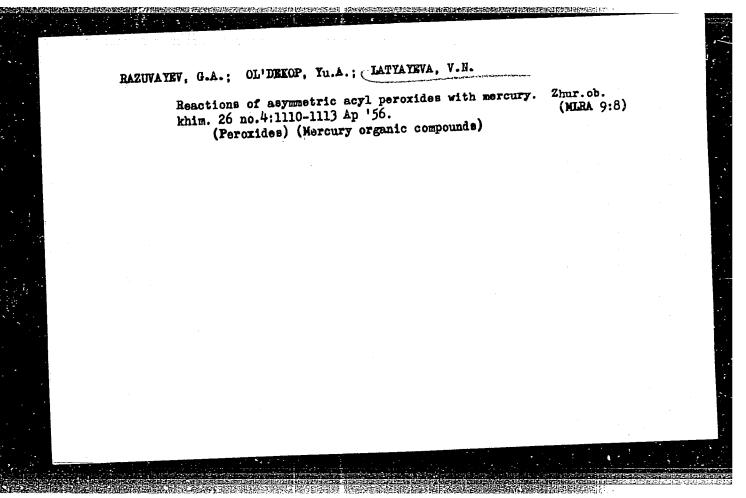
Periodical : Zhur. ob. khim. 24/2, 260-262, Feb 1954

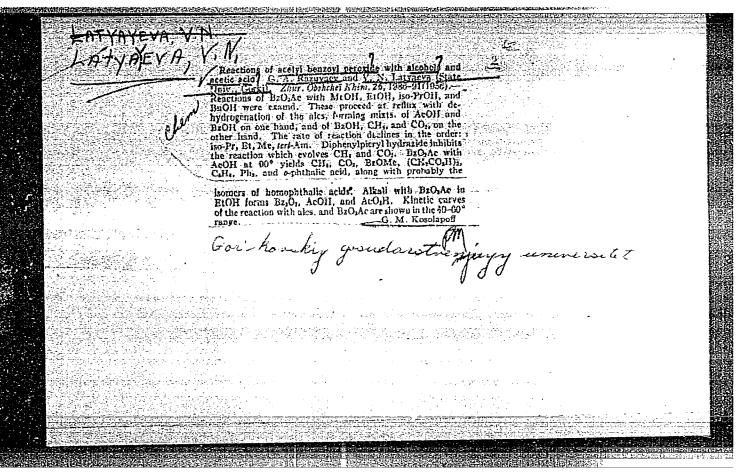
The photoreaction (exposure to ultraviolet light) of beta-mercuribispropionic Abstract

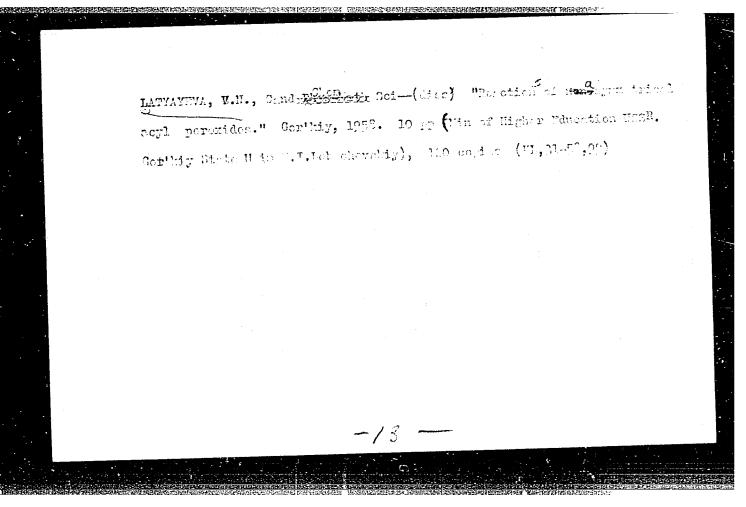
acid in solutions of methanol and monoethyl ether of ethylene glycol was investigated. The photoreaction was concluded with the separation of the mercury and formation of propionic and adipic acids. The separation of the hydro gen from the solvent by the carboxyethyl radicals was found to be instrumental in the formation of the propionic acid and the formation of adipic acid is due to the dimerization reaction of above mentioned radicals. Aldehydes were discovered in both cases. Four references: 3-USSR and 1-German (1907-1952).

State University, Gorkiy Institution:

June 19, 1953 Submitted







AUTHORS:

Razuvayev, G. A., Latyayeva, V. N.

SOV/79-28-8-48/66

TITLE:

Reactions of the Acyl Peroxides With Metals (Reaktsii atsil'nykh

perekisey s metallami)

PERIODICAL:

Zhurnal obshchey khimii, 1958, Vol. 28, Nr 8,

pp. 2233 - 2239 (USSR)

ABSTRACT:

The authors were interested in investigating more closely

the influence of different metals on the decomposition

of symmetrical and unsymmetrical acyl peroxides in solutions. Special attention was paid to the reactions of the radioactive hydrogen atom which gives up its electron pair in alcohols. For this purpose decomposition reactions of benzoyl peroxide were carried out in methyl, ethyl, and isopropyl alcohols, and in chloroform, CCl and C6H in the presence of metallic Na, Zn, Cu, Fe, Ni, Ag and Pt and at room temperature. A few reactions with phenacyl, phenacylbenzoyl, and p-nitrobenzoyl peroxide were also carried out. The experiments with benzoyl

peroxide showed that in benzene and alcohol, solutions and in the presence of Na (2% amalgam) and Zn, salts of benzoic

Card 1/3

acid form quantitatively, but no salts form from the mercury

Reactions of the Acyl Peroxides With Metals

sov/79-28-8-48/66

in the sodium amalgam. In the presence of Ni and Fe in alcohol only some of the benzoyl radicals become anions. Free benzoic acid and aldehydes were found in the reaction products in addition to the salts. The mechanism of the products in addition are probably thus: in one case the described reaction are probably thus: in one case the peroxide accepts two electrons from the metal and forms peroxide accepts two electrons from the metal and forms benzoate anions:  $(C_6H_5CO_2)_2+2n$   $\longrightarrow 2C_6H_5COO^2+2n^2+1$ . In the

other case the electron transfer occurs with the formation of a benzoate anion and a benzoyl-oxy radical (see the second reaction diagram). The latter reacts with the solvent, the alcohol. In the case of acetylbenzoyl peroxide similar the alcohol. In the case of acetylbenzoyl peroxide similar results were obtained, i.e., acetate and benzoate in the presence of Na, Hg, and Zn reacted in benzene in the same presence of Na, Hg, and Zn reacted in benzene in the same cu, and Ag acetates of the metals and free benzoic acid were found. In chloro-organic solvents chlorides of the metals were observed to form initially. In the presence of platinum a hydrogenation of the peroxide with the alcohol hydrogen occurred. The table indicates the reactions of the acyl peroxides with the metals. There are 1 figure, 1 table, and 11 refer-

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Reactions of the Acyl Peroxides With Metals

sov/79-28-8-48/66

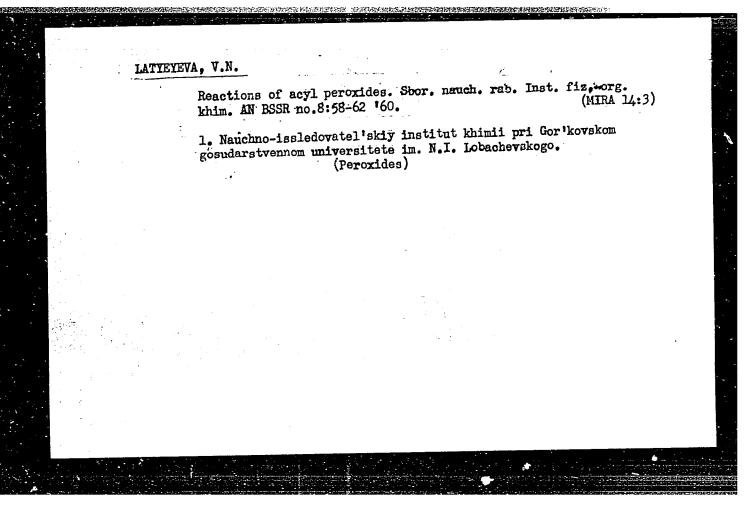
ences, 7 of which are Soviet.

ASSOCIATION: Gor'kovskiy gosudarstvennyy universitet (Gor'kiy State Uni -

versity)

SUBMITTED: July 8, 1957

Card 3/3



RAZUVAYEV, Grigoriy Alekseyevich, laureat Leninskoy premii; LATYAYEVA,
Viktoriya Nikolayevna, kand.khim.nauk; VAYNBOYM, I.B., red.;
ATROSHCHENKO, L.Ye., tekhn.red.

[Free radicals in chemistry] Svobodnye radikaly v khimii.
Moskva, Izd-vo "Znanie," 1960. 39 p. (Vsesoiuznoe obshchestvo
po rasprostraneniiu politicheskikh i nauchnykh znanii. Ser.9,
Fizika i khimiia, no.23). (MIRA 14:1)

1. Chlen-korrespondent AN SSSR (for Rezuveyev).
(Radicals (Chemistry))

s/020/60/134/003/014/020 B016/B054

5.3700 AUTHORS:

Razuvayev, G. A., Corresponding Member AS USSR,

Latyayeva, V. N., and Vyshinskaya, L. I.

TITLE:

Some Reactions of Bis-cyclopentadienyl-diphenyl Titanium

PERIODICAL:

Doklady Akademii nauk SSSR, 1960, Vol. 134, No. 3,

pp. 612-614

TEXT: The authors compare some chemical properties of  $(C_5H_5)_2^{TiAr}_2$  with those of other organometallic compounds (Ar = aryl radical). To compare reactivity, they applied the exchange reaction radical - halogen for  $(C_5H_5)_2^{TiCl}_2$  and  $(C_6H_5)_2^{Hg}$  on the one hand, and for  $(C_5H_5)_2^{Ti(C_6H_5)}_2^{Ti(C_6H_5)}_2$  and  $(C_6H_5)_2^{Hg}$  on the other. From a boiling solution of the components in

benzene or methylene chloride, they isolated a) about 20% of the expected phenyl mercury chloride from benzene, and b) nearly the theoretical yield from methylene chloride. The reaction with sublimate was carried out in CCl<sub>4</sub> or in benzene at 80°C. The main products obtained were: bis-

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Some Reactions of Bis-cyclopentadienyl-diphenyl Titanium

S/020/60/134/003/014/020 B016/B054

cyclopentadienyl-titanium dichloride and phenyl mercury chloride (1:2). The authors conclude from this ratio that in CCl<sub>4</sub> mainly (at about 70%) an exchange reaction takes place between bis-cyclopentadienyl-diphenyl titanium and the sublimate according to equation (2). In benzene solutions, the bis-cyclopentadienyl-titanium dichloride yield decreased to 24% while up to 90% of phenyl mercury chloride was formed. Further, chloro benzene, diphenyl, and calomel were isolated from the CCl<sub>4</sub> medium. Phenol also formed in the presence of atmospheric oxygen. The formation of these by-products is explained by a parallel reaction of the initial organotitanium compound with the solvent. For this reason, the authors carried out the dissociation reactions of (C<sub>5</sub>H<sub>5</sub>)<sub>2</sub>Ti(C<sub>6</sub>H<sub>5</sub>)<sub>2</sub> in different solvents.

With the exclusion of air, the original yellow color of the solution changed to dark green due to heating. The latter color corresponds to the paramagnetic form of bis-cyclopentadiene titanium (Ref. 4). The formation of chloro benzene and small amounts of diphenyl in a CCl, medium is known (Ref. 5). The authors assumed an original homolysis of the Ti—C<sub>6</sub>H<sub>5</sub> bond and the formation of a free phenyl radical; to check this assumption they allowed  $(C_5H_5)_2\text{Ti}(C_6H_5)_2$  to react with methyl- and isopropyl alcohol,

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Some Reactions of Bis-cyclopentadienyl-diphenyl

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as well as with chloroform. On the basis of the results, the authors assume the following reaction mechanism: the initial titanium compound decomposes when heated or subjected to ultraviolet radiation, along with the separation of the phenyl radical and the formation of paramagnetic, dark-green bis-cyclopentadiene titanium. The behavior of the resulting phenyl radicals depends on the type of solvent: in benzene, they yield diphenyl, whereas in alcohol solutions or in chloroform they attract the hydrogen to form benzene. All reactions mentioned remind one very much of the thermo- and photoreactions of diphenyl mercury with alcohols, with CCl4, and with chloroform, which proceed according to a free-radical of mechanism. There are 5 references: 1 Soviet and 1 US.

ASSOCIATION: Nauchno-issledovatel'skiy institut khimii Gor'kovskogo gosudarstvennogo universiteta im. N.I. Lobachevskogo (Scientific Research Institute of Chemistry of the Gor'kiy State University imeni N. I. Lobachevskiy)

SUBMITTED:

June 16, 1960

Card 3/3

S/030/61/000/004/013/015 B105/B206

AUTHORS:

Razuvayev, G. A., Corresponding Member AS USSR, Latyayeva, V. N., Candidate of Chemical Sciences, Brilkina, T. G.,

Candidate of Chemical Sciences

TITLE:

Homolytic reactions in the liquid phase

PERIODICAL: Vestnik Akademii nauk SSSR, no. 4, 1961, 124-127

TEXT: The first simpozium po gomoliticheskim reaktsiam v zhidkoy faze (Symposium on Homolytic Reactions in the Liquid Phase) held in Gor'kiy and Dzerzhinsk from December 7-10, 1960, is described. The Symposium which was attended by about 500 chemists, was convened by the Nauchnyy sovet po teorii khimicheskogo stroyeniya, kinetike i reaktsionnoy sposobnosti Otdeleniya khimicheskikh nauk Akademii nauk SSSR (Scientific Council for the Theory of Chemical Structure, Kinetics and Reactivity of the Department of Chemical Sciences AS USSR), the Gor'kovskiy nauchno-issledovatel'skiy institut khimii (Gor'kiy Scientific Research Institute of Chemistry) and the oblastnoye otdeleniye Vsesoyuznogo khimicheskogo obshchestva im. D. I. Mendeleyeva (Rayon Department of the All-Union Chemical Society imeni D. I. Men-

Card 1/5

Homolytic reactions ...

S/030/61/000/004/013/015 B105/B206

deleyev). The following reports are mentioned: By the method of electronic paramagnetic resonance, V. V. Voyevodskiy clarified the structure of benic paramagnetic resonance, v. v. voyevouskly similar radicals, and establishzene chromate cations as well as the aromatic ionic radicals, and established the formation of hydrogen atoms during the irradiation of the system Fe +H2SO4+H2O at 77°K by means of ultraviolet light; M. B. Neyman, A. L. Buchachenko reported on the formation of stable radicals which can serve as basis for the determination of active, short-lived radicals; A.N. Terenin, B.L.Kurbatov, R.F. Vasil'yev, A.A. Vichutinskiy, Q.N. Karpukhin, L.M. Postnikov, and V.Ya. Shlyapintokh reported on the method of chemiluminescence; K.S. Bagdasark yan, R. I. Milyutinskaya, E.A. Trosman, and V. A. Borovkova investigated the reactions of the phenyl- and nitrophenyl radicals with aromatic compounds by the kinetic method; V. F. Tsepalov found an expression for the rate of consumption of an arbitrary component as function of the concentration of recoing substances; N. M. Emanuel' discovered the dependence of the oxidizing of liquefied hydrocarbon on the concentration of the solvent; N. M. Emanuel', E. K. Mayzus, and I. P. Skibida reported on the production of alcohols and ketones according to the chain- and volecular method of the oxidation of n-decane; B. V. Yerofeyev reported on complementing the previous theory of primary initiating by a secondary initiating; K. I. Ivanov and Ye. D. Card 2/5

Homolytic reactions ...

S/030/61/000/004/013/015 B105/B206

FAR HERBERGER LAND BESTER HERBERGER WALLE

Vilyanskaya showed that aniline added to an oil already in a state of oxidation is converted into a product behaving similar to a peroxide radical which accelerates the reaction; B. A. Redoshkin and V. A. Shushunov showed the dual effect of metal salts of variable valency; A T Buchachenko, M. P Neyman, and K. Ya. Kaganskaya determined the average intetime or peroxid. radicals of trimethyl heptane (3.5 sec); I. V. Berezin, K. Vatsek, Go Chu, and N. F. Kazanskaya classified a number of free radicals according to their kinetic indices; Ye. N. Gur'yanova, I. G. Chernomorskaya, and M. S. Fel'dshteyn discovered the direct dependence between exchangeability of the compounds S-S, S-N, S-C and their vulcanizing activity; G. A. Razuvayev, G. G. Petukhov, Ye. V. Mitrofanova, and V. N. Istvaveva showed that the use of isotope methods permits the discovery of new reactions during the oxidation of organometallic compounds, which cannot be determined by other methods; V. A. Shushunov, Yu. A. Aleksandrov, and T. G. Brilkina submitted a scheme of the oxidation process of the organometallic compounds investigated, N. S. Vyazankin, G. A. Razuvayev, Yu. I. Dergunov, and O. A. Shchepetkova reported on the homolytic cleavage of elementary compounds; Yu. A. Ol'dekop and N. M. Mayer reported on the mechanism of the homolytic synthesis of organometallic compounds; N. P. Khyrak and V. A. Pal'm reported on the homo-Card 3/5

s/030/61/000/004/013/015 B105/B206

Homolytic reactions ...

lytic character of the formation of organomagnesium compounds: A. V. Savitskiy and Ya. K. Syrkin reported on the spectrophotometric investigations which were utilized for determining the thermodynamic indices of the oxidation reactions of ferrocene and rutheniumcene by means of iodine; G. I. Nikishin and V. D. Vorob'yev reported on the linkage of the alcohols C5-C10 to  $\alpha$ -olefins of the composition  $C_6-C_{13}$ ; G. I. Nikishin, Yu. N. Ogibin, and A. D. Petrov reported on esters of dicarboxylic acids which are linked to  $\gamma$ -olefins under formation of esters of  $\alpha$ -alkyl carboxylic acids; G. A. Razuvayev and L. S. Boguslavskaya reported on the production of glycol esters; M. G. Gonikberg and V. M. Zhulin reported on the production of an unstable polymer at a pressure of 5000 kg/cm2, which is depolymerized at customary pressure; A. P. Meshcheryakov and I. Ye. Dolgiy reported on the production of substituted cyclopropane derivatives by addition of methylene radical and its derivatives on alkene; A. N. Nesmeyanov, R. Kh. Freydlina, V. N. Kost. M. Ya. Khorlina, T. T. Sidorova, R. G. Petrova, and A. B. Terent'yev arranged the investigated radicals according to their relative stability; M. F. Shostakovskiy, Ye. N. Prilezhayeva, and L. V. Tsymbal reported on heterolytic reactions of the additions which are strictly subordinated to the rule of transaddition; G. M. Strongin reported on the conforma-

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S/030/61/000/004/013/015
Homolytic reactions ... B105/B206
tion of products of the homolytic addition of chloring on benzeng. The

tion of products of the homolytic addition of chlorine on benzene. The delegates of the Symposium expressed the wish to discuss regularly chemical problems connected with the homolytic reaction in the liquid phase.

Card 5/5

S/079/61/031/001/023/025 B001/B066

AUTHORS:

Razuvayev, G. A., Latyayeva, V. N., and Petukhov, G. G.

TITLE:

Decomposition of Acyl Peroxides in Acid Medium

PERIODICAL:

Zhurnal obshchey khimii, 1961, Vol. 31, No. 1, pp. 268 - 274

TEXT: Refs. 1 and 2 indicate the possibility of a regenerative exchange in could not be confirmed experimentally. On the basis of Refs. 1 - 5, the authors tried once more to establish the so-called "relay-transfer" of the acyloxy radicals (1) in carboxylic acids. For this purpose, the reaction of benzoyloxy- and m-nitro-benzoyloxy radicals which are more stable than the acetyloxy radicals was carried out in acetic and benzoic acid C14-labeled in the carboxyl. The separation of labeled C140, may indicate the occurrence of such an exchange, provided that the initial acids and the resultant products are stable to CO2 separation during the course of reaction. A spontaneous decarboxylation of acetic and benzoic acid at 100°C is im-Card 1/3

Decomposition of Acyl Peroxides in Acid Medium

S/079/61/031/001/023/025 B001/B066

possible. Apart from the papers of Refs. 6 - 8, no papers have been published so far on the reaction products of benzoyl and acetyl-benzoyl peroxide in benzoic acid, and of m-nitro-benzoyl peroxide in acetic acid. The authors therefore first determined the principal products of these reactions. They studied the decomposition of benzoyl-, acetyl-benzoyland m-nitro-benzoyl peroxides in acetic and benzoic acid labeled with C14 in the carboxyl. The separating carbon dioxide gas was found to contain 2 - 25 % of radioactive  $C^{14}O_2$ . Decomposition of benzoyl- and m-nitrobenzoyl peroxide in deuterated acetic acid (CH3COOD) disclosed that the R-radical of the peroxide splits off 1 - 3 % deuterium from the carboxyl group of the acid (R =  ${}^{\circ}_{6}{}^{H}_{5}$ ,  ${}^{\circ}_{6}{}^{H}_{4}{}^{NO}_{2}$ ). The separation of the labeled C140, is explained by the "relay-transfer" of the acyloxy radicals in carboxylic acid medium. The only possible source of the  ${\rm C}^{14}{\rm O}_2$  formation is thus the decarboxylation of the RC 1400° radicals of the labeled solvent. Contrary to the remaining peroxides, acetyl benzoyl peroxide gives a considerable quantity of methyl benzoate on decomposition into acids, Card 2/3

Decomposition of Acyl Peroxides in Acid Medium

S/079/61/031/001/023/025 B001/B066

especially in the case of benzoic acid (70 %), which may be explained by the reaction  $C_6H_5COOCCCH_3 \longrightarrow C_6H_5COOCH_3 + CO_2$  (2). Table 2 summarizes the experimental results on the decomposition of benzoyl-m-nitro-benzoyl peroxide dissolved in CH3COOD; they show that in the case of the phenyl radical, the deuterium separation is 3 %, and in the case of the nitrophenyl radical 1.2 %. With m-nitro-benzoyl peroxide the ratio of C6H4DNO2: C14O2 is only 1 %. S. F. Zhil'tsov is thanked for carrying out the radiometric determinations. There are 2 tables and 11 references: 5 Soviet, 4 US, and 2 British.

ASSOCIATION: Nauchno-issledovatel'skiy institut khimii pri Gor'kovskom gosudarstvennom universitete (Scientific Research Institute

of Chemistry at the Gor'kiy State University)

SUBMITTED:

January 29, 1960

Card 3/3

CIA-RDP86-00513R000928810007-6" **APPROVED FOR RELEASE: 06/20/2000** 

RAZUVAYEV, G.A.; LATYAYEVA, V.N., kand.khim.nauk; BRILKINA, T.G., kand.khim.

Homolytic reactions in the liquid phase. Vest. AN SSSR 31 no.4:124-127 Ap '61. (MIRA 14:4)

1. Chlen-korrespondent AN SSSR (for Razuvayev). (Chemical reactions)

LATYAYEVA, V.N.

5.3700

AUTHORS:

25319

S/020/61/138/005/019/025 B103/B220

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Razuvayev, G. A., Corresponding Member AS USSR, Latyaeva, V.N.,

and Vyshinskaya, L. I.

TITLE:

Reaction of benzoyl peroxide with titanocene derivatives

PERIODICAL:

Akademiya nauk SSSR. Doklady, v. 138, no. 5, 1961, 1126-1129

TEXT: The authors studied the interaction of biscyclopentadienyl titanium (C2H5)2Ti with benzoyl peroxide, since the acyl peroxides are donors of

acyloxy radicals and easily break the 0-0 bond. According to a previous paper by the authors (Ref. 1: DAN, 134, 612 (1960)),  $(C_2H_5)_2$ Ti forms on

thermal decomposition of biscyclopentadienyl-phenyl titanium in alcohol or benzene solution, is very reactive and sensitive to atmospheric oxygen. Benzoyl peroxide is known to destroy sandwich compounds completely (Posakker, Ref. 2: RZhKhim, 1959, No. 22, 78502). In the present case, the titanocene group was not decomposed in benzene or isopropyl alcohol in the cold by the action of benzoyl peroxide. The color of the solution changed instantaneously from dark green to dazzling yellow. CO<sub>2</sub> was not

X

Card 1/4

Reaction of benzoyl peroxide with...

S/020/61/138/005/019/025 B103/B220

**有我们们是这种严格的影响的影响的影响和自己的影响。** 

liberated in the reaction. The yellow crystalline product obtained in a dry nitrogen atmosphere was the expected biscyclopentadienyl titanium dibenzoate: (C5H5)2Ti(OCOC6H5)2. Since it had not yet been described, the authors also synthesized it from titanocene dichloride and silver benzoate. They checked its identity by ultimate analysis (for which T. V. Guseva is thanked), by determination of the molecular weight, the melting point, and the content of benzoate groups. (C5H5)2T1(0C0C6H5)2 can be hydrolyzed very easily, whereby the molecule of the titanocene salt decomposes and cyclopentadiene, the salt of benzoic acid, and titanic acid are formed. On alcoholysis in absolute isopropyl alcohol, cyclopentadiene, acetone, and benzoic acid were found among the reaction products. By the action of moist air, the titanocene dibenzoate molecule loses two moles of cyclopentadiene and can be converted to dibenzoxy titanium oxide  $0 = Ti(0COC_6H_5)_2$ . This product is infusible. An analogous representative of compounds of the type  $(C_5H_5)_2T1(OCOR)_2$  was obtained by the reaction of titanocene dichloride with silver acetate:  $(c_5H_5)_2$ Ti $(0000H_3)$ . It is yellow, melts at 127-130°C, and corresponds to biscyclopentadienyl

Reaction of benzoyl peroxide with ...

\$/020/61/138/005/019/025 B103/B220

titanium diacetate. On alcoholysis of the latter in absolute isopropyl alcohol, acetate groups were split off and cyclopentadiene was formed to some extent. A yellow substance was precipitated, which is insoluble in organic solvents and has a structure unknown so far. Dissolved acetone was found in the isopropyl alcohol. The yellow substance mentioned was hydrolyzed completely in dilute alkali, whereby cyclopentadiene as well as acetic and titanic acids were formed. The formation of dibenzoate indicates that, unlike ferrocene, the structure of titanocene remains unchanged in this case. The authors studied the action of benzoyl peroxide on the cyclopentadienyl compounds of tetravalent titanium, i.e., on diphenyl biscyclopentadienyl titanium. Even at room temperature, the phenyl radicals in isopropyl alcohol are replaced by the acyloxy groups of the peroxide:  $(c_5H_5)_2\text{Ti}(c_6H_5)_2 + (c_6H_5\text{COO})_2 + \text{CH}_3\text{CH}(\text{OH})\text{CH}_3 \longrightarrow (c_5H_5)\text{Ti}(\text{OCOC}_6H_5)_2$ + 206H6 + CH3COCH Thereby, biscyclopentadienyl titanium dibenzoate is formed. The phericals are converted into benzene by dehydration of the alcohol to acetone. The following absorption bands (in  $cm^{-1}$ ) were found by comparing the infrared spectra of the final and the initial compounds: (C5H5)2Ti(C6H5)2 448, 459, 606, 690, 720, 770, 822, 886, 930,

Card 3/4

Reaction of benzoyl peroxide with 25319

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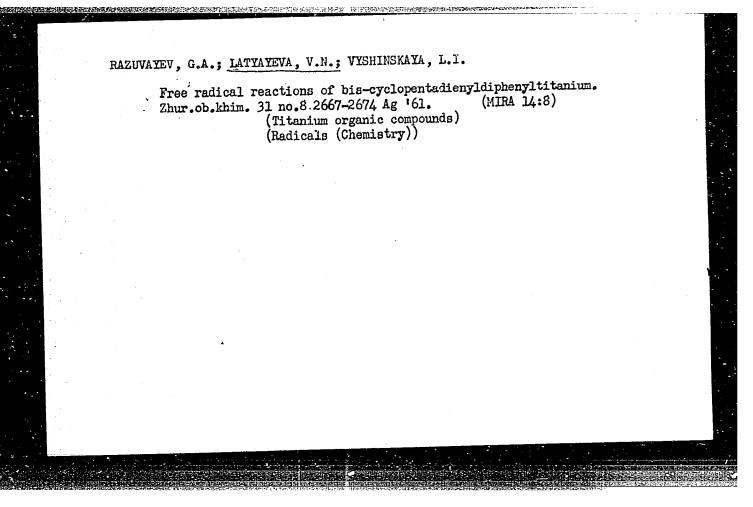
1024, 1076, 1286; (C5H5)2TiCl2: 769, 814, 828, 872, 880, 930, 1018:  $H_3$ )<sub>2</sub> 404, 520, 600, 624, 822, 865, 1024;  $(c_5H_5)_2Ti(0c0c_6H_5)_2$ 720, 83 24, 1068, 1132. The bands 822-830 and 1018-1024 cm<sup>-1</sup> are to be for the cyclopentadienyl ring. The band 865 cm<sup>-1</sup> is absent in the spectra of the initial compounds, and is interpreted as belonging to the vibrations of the Ti-O bond. There are 1 table and 3 references: 2 Soviet-bloc and 1 non-Soviet-bloc. The reference to the English-lan-guage publication reads as follows: J. D. Varma, R. C. Mehrotra (Ref. 3: Jaract. Chem. 8, 64 (1959)).

ASSOCIATION: Nauchno-issledovatel skiy institut khimii pri Gor kovskom gosudarstvennom universitete im. N. I. Lobachevskogo (Scientific Research Institute of Chemistry at Gor'kiy State University imeni N. I. Lobachevskiy)

SUBMITTED:

February 20, 1961

Card 4/4



LATYATEVA, V. N.; MALYSHEVA, A. V.; RAZUVAYEV, G. A.

Preparation of methylmercury salts. Zhur. VKHO 7 no.5:594 '62.
(MIRA 15:10)

1. Gor'kovskiy gosudarstvennyy universitet.
(Mercury compounds)

L 10285-63

Eup(1)/Epp(c)/Eut(b)/EDS\_AED\_Pc-4/Pr-4-RH/WW/HAY

ACCESSION NR: AP3000751

8/0020/63/150/003/0566/0569

AUTHOR: Razuvayev, G. A. (Corr. member AN SSSR); Latyayeva, V. N.; Maly\*sheva, A. V.; Kilyakova, G. A.

TITLE: New phenyl derivatives of Ti

SOURCE: AN SSSR. Doklady, v. 150, no. 3, 1963, 566-569

TOPIC TAGS: phenyl derivatives of Ti, PhTiCl sub 3 and Ph sub 2 Ti formation, decomposition of PhTiCl sub 3, decomposition of Ph sub 4 Ti, thermal stability of Ph sub 2 Ti

ABSTRACT: Phenyl derivatives of Ti have been synthesized for the first time by maintaining the reaction shown in formula (1) of Enclosure at approximately 90C. Of the Ti derivatives, only Ph sub 2 Ti, the first covalent metalloorganic compound of divalent Ti, was isolated in pure form. The formation of PhTiCl sub 3 (I) was confirmed by the following reactions: 1) the reaction shown in formula (2) of Enclosure; 2) decomposition of I to form diphenyl and TiCl sub 3; and 3) decomposition of I in C sub 14-tagged benzene to diphenyl

Cord 1/32

L 10285-63

ACCESSION NR: AP3000751

0

containing no C sup 14. The formation of diphenyl prompted the study of reactions of TiCl sub 4 with varying amounts of Ph sub 2 Hg or PhLi in tetrahydrofuran. Better results were obtained with PhLi. An intense black discoloration was observed at room temperature when the TiCl sub 4/PhLi ratio was 4/1. At =70C thermally unstable orange=red crystals were formed. The assumption that the latter were Ph sub 4 Ti (II) which could not be isolated was confirmed by reaction with HgCl sub 2 as shown in formula (3) of Enclosure. In the formation of II, a black substance was isolated which, after recrystallization in saturated hydrocarbons (n=nonane), formed a black crystalline compound which ignites spontaneously in air. The compound proved to be diphenyl titanium (III) formed by the decomposition of II as shown in formula (4) of Enclosure. Compound III is stable but extremely 0 sub 2-sensitive and decomposes slowly in a sealed ampoule at 200C into diphenyl and metallic titanium mirror. The composition of III was confirmed by chemical analysis and by its reactions. Whether the structure of III is monomeric or polymeric was not determined. Orig. art. has: 6 formulas.

ABSOCIATION: none

SUBMITTED: 16Feb53 SUB CODE: 00 DATE ACQ: 21Jun63 NO REF SOV: CO1 ENCL: 01 OTHER: 005

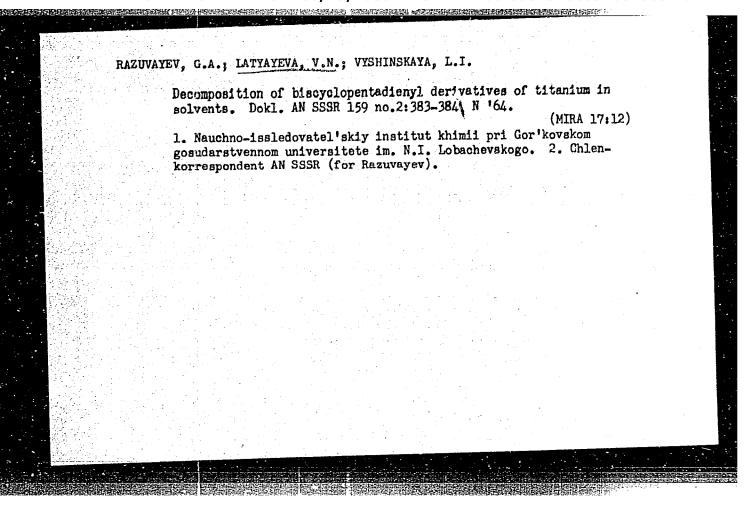
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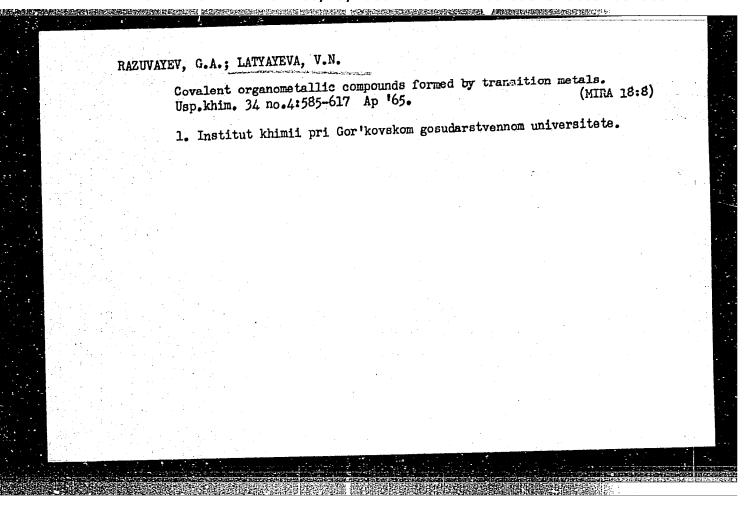
1 24832-65 EWT(m)/EPF(c)/EPR/EWP(j) Pc-li/Pr-li/Pa-li ACCESSION NR: AP4049488 8/0020/64/159/002/0383/0384 AUTHOR: Razuvayev, G.A., (Corresponding member AN SSSR), Latyayeva, V.N., Vy\*shinskaya, L.I. TITLE: Decomposition of biscyclopentadienylderivatives of titanium in solvents SOURCE: AN ESSR. Doklady\*, v. 159, no. 2, 1964, 383-384 TOPIC TAGS: biscyclopentadienyl titanium, organotitanium compound, electron paramagnetic resonance, alkyltitanium solvation ABSTRACT: The purpose of the work was to determine if there is any difference in the thermal decomposition of biscyclopenta-dienyldiethyl titanium in n-hexane and tetrahydrofuran. Studies of EPR spectra indicated that decomposition takes place by successive rupture of radicals with formation of intermediate compounds of trivalent Ti, which further decompose to compounds of divalent Ti that give no EPR signal. Both solvents behave identically. Thus, it can be assumed that the process takes place in accordance with (1) (C4H1)1TIR3 → (C4H1)3TIR + R (2)(CHI) TIR - (CHI) TI+ R" Card

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ACCESSION NR: AP40494	38	1
The nature of the radical distribution reactions of $(C_5H_5)_2$ 3	oes not affect the general nature TR2: Orig. art. has: 1 figure a	of the exchange and decomposing ocheracial equations.
ASSOCIATION: Nauchno-l nom universitete lm. N.I. Gor'kiy State University)	ssledovateľskiy instibut khimii p Lobachevskogo (Sc <u>ientific Reses</u>	rl Gor'kovskom gosudarstven- rch Institute of Chemistry,
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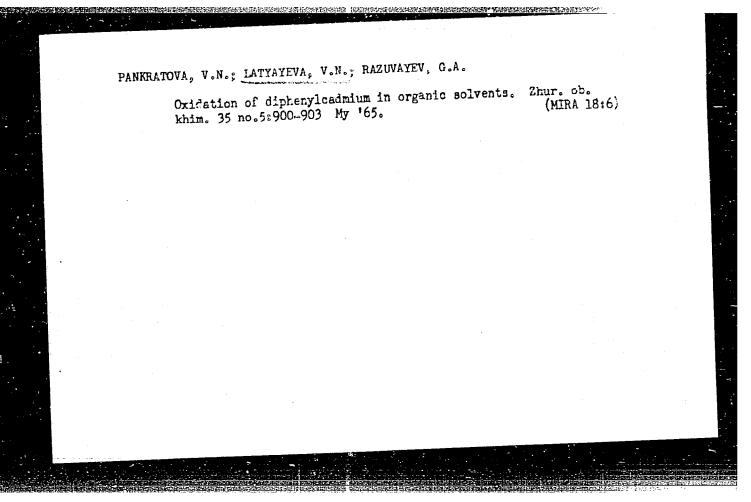
ACCESSION NR: APA040951  (Corresponding member AN SSSR)  AUTHOR: Razuvayev, G. A., Latyayeva, V. N.; Vy*shinskaya, L. I.; Vy*shinskiy, N. N.  TITLE: New monocyclopentadienyl derivatives of titanium  SOURCE: AN SSSR. Doklady*, v. 156, no. 5, 1964, 1121-1123  TOPIC TAGS: titanium, titanium derivative, monocyclopentadienyl derivative, Ti monocyclopentadienyl derivative, phenol, cyclopentadienyl timethyltitane, dipenyl mercury, phenyl mercury chloride, organotitanium compound  ABSTRACT: The authors analyzed reactions wherein the Cl atoms in monocyclopentadienyl titanium trichloride were replaced with phenyl groups. G. A. Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction of titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals. of titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals. The authors therefore initially analyzed the exchange reaction of diphenyl mercury with C. H. Ti Cl., at a 3 to 1 ratio in a benzene solution at room temperature. The following new compounds were identified:    CH.TICL, + 3 (C.H.).H. → C.H.TI (C.H.).H. (C.H.).TI (C.H.)	4	the same and the s
SOURCE: AN SSSR. Doklady*, v. 156, no. 5, 1964, 1121-1123  TOPIC TAGS: titanium, titanium derivative, monocyclopentadienyl derivative, Ti monocyclopentadienyl derivative, phenol, cyclopentadienyl timethyltitane, dipenyl mercury, phenyl mercury chloride, organotitanium compound  ABSTRACT: The authors analyzed reactions wherein the Cl atoms in monocyclopentadienyl titanium trichloride were replaced with phenyl groups. G. A. Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction of titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals. The authors therefore initially analyzed the exchange reaction of diphenyl mercury with C5 H5 Ti Cl3 at a 3 to 1 ratio in a benzene solution at room temperature. The following new compounds were identified:    CH4TICl+3 C4H5L+3 C4H5HgCl+[C4H5TICL-1]. (2)   C4H5TICl+3 C4H5L-3 C4H5HgCl+C4H5TICL-1 (2)   C4H5TICL+3 C4H5L-3 C4H5HgCl+C4H5TICL-1 (3)	" 1	DOLON ILLE
TOPIC TACS: titanium, titanium derivative, monocyclopentadienyl derivative, Ti monocyclopentadienyl derivative, phenol, cyclopentadienyl timethyltitane, dipenyl mercury, phenyl mercury chloride, organotitanium compound  ABSTRACT: The authors analyzed reactions wherein the Cl atoms in monocyclopentadienyl titanium trichloride were replaced with phenyl groups. G. A. Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction of titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals. The authors therefore initially analyzed the exchange reaction of diphenyl mercury with C5 H5 Ti Cl3 at a 3 to 1 ratio in a benzene solution at room mercury with C5 H5 Ti Cl3 at a 3 to 1 ratio in a benzene solution at room temperature. The following new compounds were identified:    C4H5TICl3+3C4H5LI→C4H5TI(C4H3)3+3LICL. (2)   C4H5TICl3+3C4H5LI→C4H5TI(C4H3)3+3LICL. (3)	L -	· · · · · · · · · · · · · · · · · · ·
monocyclopentadienyl titanium trichloride were replaced with photoside in the reaction Razuvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction of titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals. The authors therefore initially analyzed the exchange reaction of diphenyl mercury with C <sub>5</sub> H <sub>5</sub> Ti Cl <sub>2</sub> at a 3 to 1 ratio in a benzene solution at room temperature. The following new compounds were identified:    C <sub>4</sub> H <sub>5</sub> TiCl <sub>2</sub> + 3 (C <sub>4</sub> H <sub>5</sub> ) <sub>4</sub> H <sub>5</sub> - 3C <sub>4</sub> H <sub>5</sub> H <sub>5</sub> Cl + [C <sub>4</sub> H <sub>5</sub> Ti (C <sub>4</sub> H <sub>5</sub> ) <sub>3</sub> . (1)   C <sub>4</sub> H <sub>5</sub> TiCl <sub>3</sub> + 3C <sub>4</sub> H <sub>5</sub> Ll → C <sub>4</sub> H <sub>5</sub> Ti (C <sub>4</sub> H <sub>5</sub> ) <sub>4</sub> + 3LiCl. (2)   C <sub>4</sub> H <sub>5</sub> Ti (C <sub>4</sub> H <sub>5</sub> ) <sub>4</sub> + 3H <sub>5</sub> Cl <sub>4</sub> → 3C <sub>4</sub> H <sub>5</sub> H <sub>5</sub> Cl + C <sub>4</sub> H <sub>5</sub> TiCl <sub>5</sub> . (3)	TO Ti di	IC TAGS: titanium, titanium derivative, monocyclopentadienyl derivative, monocyclopentadienyl derivative, phenol, cyclopentadienyl timethyltitane, enyl mercury, phenyl mercury chloride, organotitanium compound
temperature. The following new compounds were identified. (a)    C <sub>6</sub> H <sub>6</sub> TiCl <sub>2</sub> + 3 (C <sub>6</sub> H <sub>8</sub> ) <sub>2</sub> Hg → 3C <sub>6</sub> H <sub>6</sub> HgCl + [C <sub>6</sub> H <sub>6</sub> Ti (C <sub>6</sub> H <sub>8</sub> ) <sub>2</sub> ]. (2)    C <sub>6</sub> H <sub>6</sub> TiCl <sub>2</sub> + 3C <sub>6</sub> H <sub>6</sub> Ll → C <sub>6</sub> H <sub>6</sub> Ti (C <sub>6</sub> H <sub>8</sub> ) <sub>2</sub> + 3LiCl. (2)    C <sub>6</sub> H <sub>6</sub> Ti (C <sub>6</sub> H <sub>8</sub> ) <sub>2</sub> + 3HgCl <sub>2</sub> → 3C <sub>6</sub> H <sub>6</sub> HgCl + C <sub>6</sub> H <sub>6</sub> TiCl <sub>2</sub> . (3)	mo Ra of Th	ocyclopentadienyl titanium trichloride were replaced with playing the reaction uvayev it Cl (DAN, 150 (1963) 566) Previously showed that, during the reaction titanium tetrachloride, all four Cl atoms are replaced by phenyl radicals.  authors therefore initially analyzed the exchange reaction of diphenyl authors therefore initially analyzed in a benzene solution at room
. Card 1/2	te	perature. The following new compounds were identified:    CaHaTICla + 3 (CaHa)aHg - 3CaHaHgCl + [CaHaTi (CaHa)a]. (2)   CaHaTiCla + 3CaHaLi - CaHaTi (CaHa)a + 3LiCl. (2)   CaHaTi (CaHa)a + 3HgCla - 3CaHaHgCl + CaHaTiCla. (3)

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	ACCESSION NR: AP4040951	
	$\begin{array}{c} C_{6}H_{5}TI\left(C_{6}H_{5}\right)_{5} + u_{30} \cdot C_{5}H_{7}OH \rightarrow 3C_{6}H_{5}TI\left(0 \cdot u_{30} \cdot C_{6}H_{7}\right)_{3}. & (4) \\ C_{6}H_{5}TI\left(0 \cdot u_{30} \cdot C_{5}H_{7}\right)_{3} + 3H\overline{CI} \rightarrow C_{6}H_{5}TI\overline{CI}_{3} + 3u_{30}\overline{C_{5}}H_{7}OH. & (5) \\ C_{6}H_{5}TI\left(\overline{C_{6}H_{5}}\right)_{2} & C_{5}H_{5}TI\overline{C_{6}H_{5}} + C_{6}H_{6} + \frac{1}{2}C_{6}H_{5} \rightarrow C_{6}H_{5}. & (6) \\ & C_{5}H_{5}TIC_{6}H_{5} \rightarrow C_{6}H_{6} + TI + [C_{6}H_{6}]. & (7) \\ & C_{6}H_{5}TIC_{6}H_{5} \rightarrow C_{6}H_{5}H_{5}CI + [C_{6}H_{5}TICI], & (8) \\ & I[C_{6}H_{5}TICI] + 2H_{5}CI_{2} \rightarrow C_{6}H_{5}TICI_{3} + H_{5}CI_{2}. & (9) \end{array}$	
	Authors conclude that the bonding of the titanium atom with the cyclopentadienyl ring in the examined compounds is very similar to a ferroone bond. Orig. art. has: 11 Formulas.	
	ASSCCIATION: Nauchno-issledovatel'skiy institut khimii pri Gor'kovskom gosudarstvennom universitete im. N. I. Lobachevskogo (Scientific Research Institute for Chemistry of Gorki State University)	
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Reactions of biscyclopentadicnyldithenyltitanica with benzyl chloride and triphenylchloromethane. Thur. ob. knim. 35 no.1: 169-174 Ja '65.



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LATYAYEVA, V.N.; RAZUVAYEV, G.A.; KILYAKOVA, G.A.

Diphenyltitanium complexes with tetrahydrofuran and ammonia. Zhur. ob. khim. 35 no.8:1498-1499 Ag '65. (MIRA 18:8)

1. Nauchno-issledovatel'skiy institut khimii pri Gor'kovskom gosudarstvennom universitet.

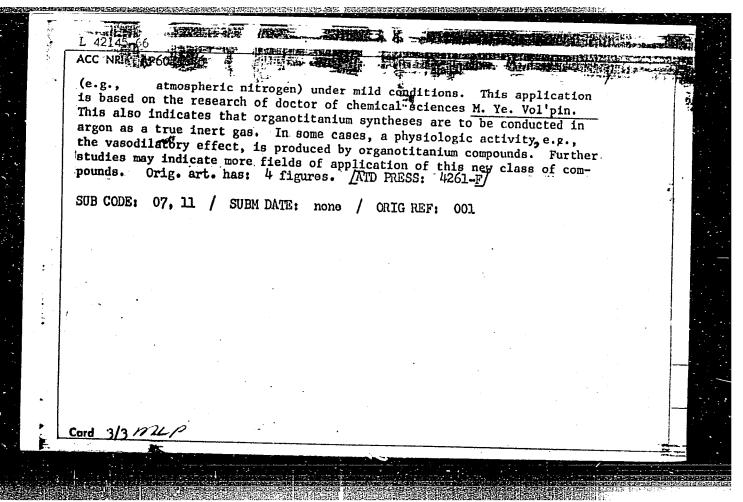
RAZUVAYEV, G.A.; MINSKER, K.S.; LATYAYEVA, V.N.; SANGALOV, Yu.A.

Polymerization of viryl chloride initiated by the reaction of carbon tetrachloride with ti' virum organometallic compounds. Dokl. AN ESSR 163 no.4:906-908 Ag \*65.

1. Mauchno-issledovatel'skiy institut khimii pri Gor'kovskom Gosudaretvennem universitete im. N.I.Lobachevskogo. 2. Chlenkorrespondent AN SECR (for Razuvayev).

WW/RM EWT(m)/EWP(j)/T IJP(c) SOURCE CODE: UR/0026/66/000/005/0048/0053 42145-66 ACC NR: AP6016846 AUTHOR: Razuvayev, G. A. (Corresponding member AN SSSR); Latyayeva, V. N. 10 (Candidate of chemical sciences) ORG: Gor'kovskiy State University im. N. I. Lobachevskiy (Gor'kovskiy gosudarstvennyy universitet) TITIE: New class of compounds. Research and discovery of organotitanium derivatives SOURCE: Priroda, no. 5, 1966, 48-53 TOPIC TAGS: titanium, organotitanium compound, metal industry, pi bonded organometallic compound, chemical bonding, chemical synthesis, free radical, polymerization reaction mechanism, argon, biochemistry ABSTRACT: This popular science type article reflects a special Soviet interest in titanium and its compounds. The summary of the article even states. that wide-spread opinion is being formed on replacing the "iron age" with the "titanium age". It is emphasized that the monument in Moscow erected to honor the conquerors of space is coated with this metal. It is noted in the article that the use of metallic titanium and its alloys is handicapped by time-consuming and costly refining. However, since titanium is a transition element it is of considerable interest not only in its metallic form, but also in its organometallic compounds. The 546.821 Card 1/3

L 42145-66 ACC NR: AP6016846 alkoxy titanium derivatives which are used for the preparation of heat resistant plastics mainly due to the research made by Academician K. A. Andrianov, are mentioned first. Secondly, the Ziegler-Natt catalysts are of It is emphasized that the organotitanium compounds supposedly formed in considerable importance. the course of the reactions promoted by these catalysts, and many otherknown compounds, e.g.. biscyclopentadienyltitanium belong to the class of the so-called "sandwich" compounds or metallocenes in which vacancies in the titanium atom shell are filled by  $\pi$ -electrons of organic radicals. However, true covalent organotitanium compounds were considered to be unattainable until attempts were made to synthesize them at very low temperatures and in an inert gas (argon) atmosphere. Thus, a new class of titanium compounds was obtained: mixed sandwich-covalent compounds and purely covalent compounds. A peculiarity of the latter is their intense color. which is contrary to the colorless covalent organic compounds of nontran-Covalent organotitanium compounds are not stable at room temperature, are easily oxidized in the air, and are hydrolyzed by moisture. In some cases, these compounds decompose according to the free radical mechanism and can inititate the polymerization of vinyl monomers. Another potential practical application of the reactivity of covalant organic compounds of titanium or some transition metals is the fixation of molecular nitrogen Card 2/3



11410-6/ E#T(in)/EMP(j) IGH SOURCE CODE: UR/0079/66/036/008/1491/1498		
AUTHOR: Razuvayev, G. A.; Latyayeva, V. N.; Vyshinskaya, L. I.; Kilyakova, G. A. ORG: Scientific Research Institute, Gor'kly State University im. N. I.  ORG: Scientific Research Institute, Gor'kly State University im. N. I.		
universitete) TITLE: Some reactions of Bis-cyclopentadienyltitanium and monocyclopentadienyl-		
phenyltitanium  SOURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1491-1498  SOURCE: Zhurnal obshchey khimii v. 36, no. 8, 1966, 1491-1498  TOPIC TAGS: organotitanium compound, thermal decomposition, chemical bonding  TOPIC TAGS: organotitanium compound, thermal reactions of decomposition of pi-  ABSTRACT: In a study of whether thermal reactions of decomposition for different R,		
ABSTRACT: In a study of whother the standard common for different type cyclopentadienyl compounds of tetravalent titanium are common for different type cyclopentadienyl derivatives cyclopentadienyl derivatives		
the known reactions of the control o		
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l lower valence, analogously to my hond was unaffected. The	,	
diphenyltitanium and diphenyl. The place of the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to thermal decomposition increased in the series: stability of the compounds to the compounds to the compounds to the compounds to the compound of the		
$(c_6H_5)_{i_1}^{Ti} < (c_5H_5)^{Ti}(c_6H_5/3)$		
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for the first time, and possessed one pi-bond C5H5-Ti and three sigma-bonds C6H5-Ti. The products of thermal decomposition: (C6H5)2Ti, C5H5TiC6H5, and (C5H5)2Ti were more stable to the action of high temperatures, but were extremely readily oxidized. The reactions of (C6H5)2Ti, C6H5TiC5H5, and (C5H5)2Ti with halo-derivatives included cleavage of the phenyltitanium bonds and their replacement by chlorine-titanium bonds. In the reaction of these compounds with chloroform, carbon tetrachloride, mercuric chloride, and hydrogen chloride, the C5H5Ti and (C5H5)2Ti groups were unaffected. The titanium-containing final products were TiCl<sub>1</sub>, C5H5TiCl<sub>3</sub>, and (C5H5)2TiCl<sub>2</sub>, respectively. The reactions of organotitanium compounds considered illustrate the relative stability of the pi-bond C5H5Ti to the action of temperatures, halo-derivatives and other reagents in comparison with the sigma-bond Ti-R. [JPRS: 38,970]

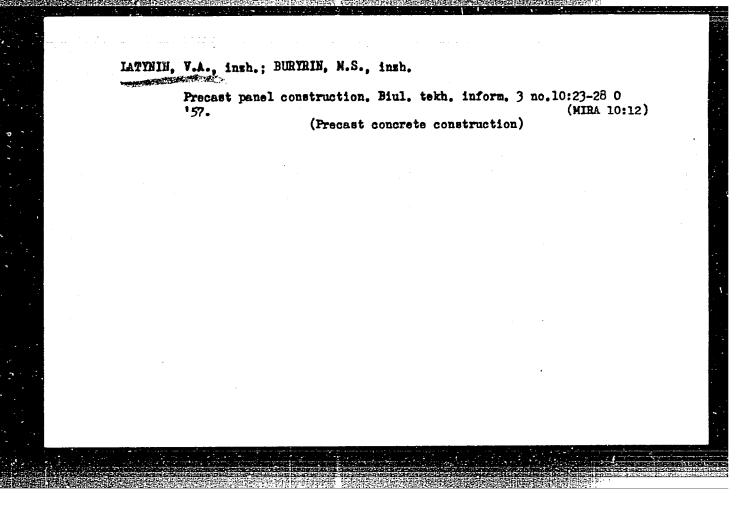
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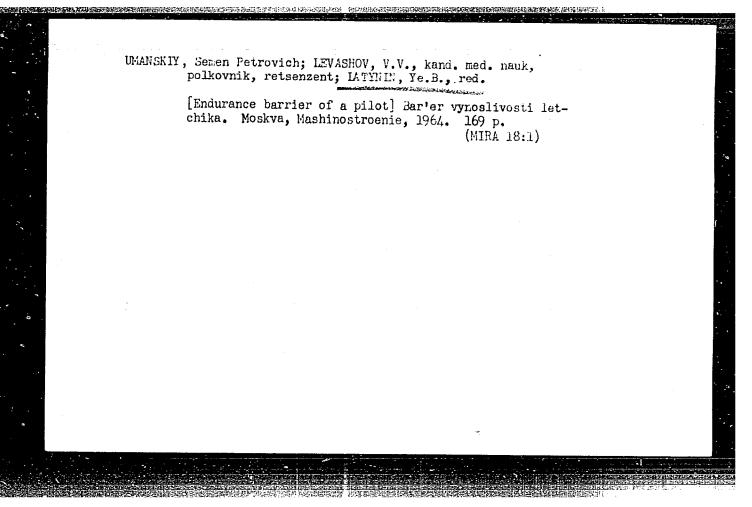
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Upper Silesian industrial area of the Polish People's Republic. Geog. v shkole 21 no. 4:16-25 Jl-Ag '58. (MIRA 11:7) (Poland--Industries)

Characteristics of the production and territorial structure of
Katowice Province. Vest. Mosk. un. Ser. 5: Geog. 19 no.1:50-55
Ja-F '64. (MIRA 17:4)

1. Kafedra ekonomicheskoy geografii Moskovskogo finansovogo
instituta.





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VASIL'YEV, G.S.; MIKIRTUMOV, E.B., kandidat tekhnicheskikh nauk, redaktor;

LATTNIN, Ye.V., redaktor; ZUDAKIN, I.M., tekhnicheskiy redaktor.

[Principles of flight applied to airplane models with flapping wings] Genovy poleta modelei s mashushchimi kryl'iami, Pod red.

E.B.Mikirtumova. Moskva, Gos. izd-vo oboronnoi promyshl., 1953.

123 p. [Microfilm] (MIRA 7:10)

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LATYMIN, Ye.V.

RESHYANSKIY, F.M.; SKLYANKIN, A.N.; USOVA, A.V.: LATTHIN, Ye.V., redaktor; Mukhina, T.N., tekhnicheskiy redaktor

[Industrial excursions in physics; a handbook for teachers]

Protzvodstvennye ekskursii po fizike; pesohie dilia uchitelei.

Moskva, Izd-vo Akad, pedagog. nauk RSFSR, 1954. 233 p. (MIRA 8:5)

(Physics-Study and teaching)(Scheol excursions)

(Technology)

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ARDAB YEVSKIY, A.I.; VOROPAYEVA, V.G.; GRINEVA, K.I.; VISHNEVSKIY, A.†a.,
inzhener, redaktor: LATVHUM, Ye.V., inzhener, xaveduyushchiy
redaktsiey; SHEKHTMAM, X.A., izdatel skiy redaktor; NOZHIW, V.P.,
tekhnicheskiy redaktor.

[Manual on calculations for super-high frequency antennas] Posobie
po raschetu antenn sverkhvysokikh chastot. Pod obshchei red, K.I.
Grinevoi. Moskva, Gos.izd-vo obor.prowyshl. 1957. 70 p.

(Antennas (Blectronics))

(Antennas (Blectronics))